

To: August Koster(August@signifyanalytics.com)
Subject: U.S. Trademark Application Serial No. 97135716 - SIGNIFY ANALYTICS
Sent: November 07, 2023 06:45:27 PM EST
Sent As: tmng.notices@uspto.gov

Attachments

[screenshot-www-bio-techne-com-reagents-cell-culture-reagents-16993949062271](#)
[screenshot-www-bio-techne-com-search-16993950303831](#)
[screenshot-www-bio-techne-com-diagnostics-quantidex-oncology-testing-kits-16993951075261](#)
[screenshot-mdmaxx-com-products-abbott-195-160-binaxnow-covid19-at-home-antigen-self-test-2-tests-16993952247991](#)
[screenshot-mdmaxx-com-products-abbott-195-160-binaxnow-covid19-at-home-antigen-self-test-2-tests-16993953013531](#)
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[Amazon Reagent kits1.jpg](#)
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United States Patent and Trademark Office (USPTO)
Office Action (Official Letter) About Applicant's Trademark Application

U.S. Application Serial No. 97135716

Mark: SIGNIFY ANALYTICS

Correspondence Address:

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COSTA MESA CA 92627
UNITED STATES

Applicant: August Koster

Reference/Docket No. N/A

Correspondence Email Address: August@signifyanalytics.com

REQUEST FOR RECONSIDERATION AFTER FINAL ACTION DENIED

Issue date: November 7, 2023

This Office action is in response to applicant's communication filed on October 17, 2023. Applicant's amendment to the identification of goods has been accepted except for the format of the wording "in Class 5," as this inclusion does not meet the standardized stylistic form used by the Office. The full identification now reads as:

International Class 005: Drug testing kits comprised of diagnostic test strip, chemical reagents and buffers, cotton swabs, test tube and test tube holder and printed instructions that test for the presence of fentanyl; Drug testing kits diagnostic test strip, chemical reagents and buffers, cotton swabs, test tube and test tube holder and printed instructions that test for the presence of drugs or alcohol; COVID-19 test kits comprised of diagnostic test strip, chemical reagents and buffers, cotton swabs, test tube and test tube holder and printed instructions; Diagnostic kits comprised of diagnostic test strip, chemical reagents and buffers, cotton swabs, test tube and test tube holder and printed instructions to test for the presence of viruses; Diagnostic kits diagnostic test strip, chemical reagents and buffers, cotton swabs, test tube and test tube holder and printed instructions to test fecal matter; Drug testing kits comprised of medical diagnostic reagents and assays for testing body fluids; Diagnostic kits consisting primarily of monoclonal antibodies, buffers, and reagents for use in disease testing; Diagnostic preparations for medical purposes; Specimen collection kits for use in virus testing comprised primarily of swabs for medical purposes, chemical reagents, buffers and assays, test tubes, and printed instructions; Diagnostic test kits to determine sperm fertility; Fertility enhancement preparations; Ovulation test kits; Medical diagnostic test strips for use in the field of reproduction; **Genetic identity tests comprised of reagents for medical purposes, all of the foregoing for home use and not for professional use.**

The wording regarding the classification appears at the beginning of the identification. All goods that follow the wording are classified in Class 5.

Applicant's request for reconsideration is denied. See 37 C.F.R. §2.63(b)(3). The trademark examining attorney has carefully reviewed applicant's request and determined the request did not: (1) raise a new issue, (2) resolve all the outstanding issue(s), (3) provide any new or compelling evidence with regard to the outstanding issue(s), or (4) present analysis and arguments that were persuasive or shed new light on the outstanding issue(s). TMEP §§715.03(a)(ii)(B), 715.04(a).

Accordingly, the following requirement(s) and/or refusal(s) made final in the Office action dated April 18, 2023 are **maintained and continued**:

- The refusal to register under Section 2(d) of the Trademark Act, 15 USC Section 1052(d), based on likelihood of confusion with U.S. Registration No. 2112978;
- The requirement for a disclaimer of "Analytics" apart from the mark.

See TMEP §§715.03(a)(ii)(B), 715.04(a).

LIKELIHOOD OF CONFUSION - REFUSAL CONTINUED TO BE MADE FINAL

Marks are compared in their entireties for similarities in appearance, sound, connotation, and commercial impression. *Stone Lion Capital Partners, LP v. Lion Capital LLP*, 746 F.3d 1317, 1321, 110 USPQ2d 1157, 1160 (Fed. Cir. 2014) (quoting *Palm Bay Imps., Inc. v. Veuve Clicquot Ponsardin Maison Fondee En 1772*, 396 F.3d 1369, 1371, 73 USPQ2d 1689, 1691 (Fed. Cir. 2005)); TMEP §1207.01(b)-(b)(v). "Similarity in any one of these elements may be sufficient to find the marks

confusingly similar.” *In re Inn at St. John’s, LLC*, 126 USPQ2d 1742, 1746 (TTAB 2018) (citing *In re Davia*, 110 USPQ2d 1810, 1812 (TTAB 2014)), *aff’d per curiam*, 777 F. App’x 516, 2019 BL 343921 (Fed. Cir. 2019); TMEP §1207.01(b).

When comparing marks, “[t]he proper test is not a side-by-side comparison of the marks, but instead whether the marks are sufficiently similar in terms of their commercial impression such that [consumers] who encounter the marks would be likely to assume a connection between the parties.” *Cai v. Diamond Hong, Inc.*, 901 F.3d 1367, 1373, 127 USPQ2d 1797, 1801 (Fed. Cir. 2018) (quoting *Coach Servs., Inc. v. Triumph Learning LLC*, 668 F.3d 1356, 1368, 101 USPQ2d 1713, 1721 (Fed. Cir. 2012)); TMEP §1207.01(b). The proper focus is on the recollection of the average purchaser, who retains a general rather than specific impression of trademarks. *In re Ox Paperboard, LLC*, 2020 USPQ2d 10878, at *4 (TTAB 2020) (citing *In re Bay State Brewing Co.*, 117 USPQ2d 1958, 1960 (TTAB 2016)); *In re Inn at St. John’s, LLC*, 126 USPQ2d 1742, 1746 (TTAB 2018); TMEP §1207.01(b); see *In re St. Helena Hosp.*, 774 F.3d 747, 750-51, 113 USPQ2d 1082, 1085 (Fed. Cir. 2014).

Consumers are generally more inclined to focus on the first word, prefix, or syllable in any trademark or service mark. See *Palm Bay Imps., Inc. v. Veuve Clicquot Ponsardin Maison Fondée En 1772*, 396 F.3d 1369, 1372, 73 USPQ2d 1689, 1692 (Fed. Cir. 2005) (finding similarity between VEUVE ROYALE and two VEUVE CLICQUOT marks in part because “VEUVE . . . remains a ‘prominent feature’ as the first word in the mark and the first word to appear on the label”); *Century 21 Real Estate Corp. v. Century Life of Am.*, 970 F.2d 874, 876, 23 USPQ2d 1698, 1700 (Fed. Cir. 1992) (finding similarity between CENTURY 21 and CENTURY LIFE OF AMERICA in part because “consumers must first notice th[e] identical lead word”); see also *In re Detroit Athletic Co.*, 903 F.3d 1297, 1303, 128 USPQ2d 1047, 1049 (Fed. Cir. 2018) (finding “the identity of the marks’ two initial words is particularly significant because consumers typically notice those words first”).

MARKS ARE CONFUSINGLY SIMILAR: In this case the applicant has argued that the term "Signify" as used by applicant and registrant have differing commercial impressions. However, the term "Signify," even when considering applicant's own dictionary definitions, has a highly limited meaning which is to be a sign of or to show by a conventional token. In short, "Signify" means something that shows or proves the existence of a factual matter. Applicant has further alleged that the addition of the term "Analytics" is distinctive and that it alters the meaning of "Signify" alone. However, the examining attorney disagrees. The term "Analytics" means the method of logical analysis. When used in connection with applicant's identified goods, which are medical test kits comprised of chemical reagents to test for the existence or presence of drugs, viruses, hormones or other substances in the body, the term merely describes the primary purpose or function of the goods, namely, that it is a tool used to provide logical analysis of a medical fact i.e., the presence of certain hormones means that the person tested is pregnant, the presence of certain viruses means the person has the flu or Covid-19, the presence of certain chemicals means that the person has cancer or is under the influence of a controlled substance or alcohol. The addition of "Analytics" does not significantly alter the commercial impression of the dominant element, SIGNIFY, which in both marks is spelled the same, sounds the same, a presents a similar overall meaning. This is not the same as the *Patron Spirits International AG. vs. Conyngham Brewing Company* case cited by applicant in its Request for Reconsideration where the mark PIRATE PISS for beer was found not to be confusingly similar to PYRAT for rum. The term "Piss" has no descriptive meaning relative to the identified goods, beer. Thus, the addition of PISS to PIRATE, does materially change the overall meaning of the first element, Pirate. The mark is now no longer just a "pirate," but excrement from a pirate. Had the marks been PIRATE and PIRATE BEER, the result may have been different precisely because the additional

generic term does not make the mark distinguishable from PIRATE alone. Disclaimed matter or matter that is descriptive of or generic for a party's goods and/or services is typically less significant or less dominant when comparing marks. See *In re Detroit Athletic Co.*, 903 F.3d 1297, 1305, 128 USPQ2d 1047, 1050 (Fed. Cir. 2018) (citing *In re Dixie Rests., Inc.*, 105 F.3d 1405, 1407, 41 USPQ2d 1531, 1533-34 (Fed. Cir. 1997)); *Made in Nature, LLC v. Pharmavite LLC*, 2022 USPQ2d 557, at *41 (TTAB 2022); TMEP §1207.01(b)(viii), (c)(ii).

GOODS ARE RELATED: The compared goods and/or services need not be identical or even competitive to find a likelihood of confusion. See *On-line Careline Inc. v. Am. Online Inc.*, 229 F.3d 1080, 1086, 56 USPQ2d 1471, 1475 (Fed. Cir. 2000); *Recot, Inc. v. Becton*, 214 F.3d 1322, 1329, 54 USPQ2d 1894, 1898 (Fed. Cir. 2000); TMEP §1207.01(a)(i). They need only be "related in some manner and/or if the circumstances surrounding their marketing are such that they could give rise to the mistaken belief that [the goods and/or services] emanate from the same source." *Coach Servs., Inc. v. Triumph Learning LLC*, 668 F.3d 1356, 1369, 101 USPQ2d 1713, 1722 (Fed. Cir. 2012) (quoting *7-Eleven Inc. v. Wechsler*, 83 USPQ2d 1715, 1724 (TTAB 2007)); TMEP §1207.01(a)(i); see *Made in Nature, LLC v. Pharmavite LLC*, 2022 USPQ2d 557, at *44 (TTAB 2022) (quoting *In re Jump Designs LLC*, 80 USPQ2d 1370, 1374 (TTAB 2006)).

Determining likelihood of confusion is based on the description of the goods and/or services stated in the application and registration at issue, **not on extrinsic evidence of actual use**. See *In re Detroit Athletic Co.*, 903 F.3d 1297, 1307, 128 USPQ2d 1047, 1052 (Fed. Cir. 2018) (citing *In re i.am.symbolic, llc*, 866 F.3d 1315, 1325, 123 USPQ2d 1744, 1749 (Fed. Cir. 2017)).

Applicant has included in the record printouts from the registrant's website and most recent renewal application and argues that the evidence shows that the registrant's goods are only sold to medical professionals and not to home users as applicant has since amended its application. However, the registrant's identification is not restricted to professional or medical provider use only and cannot be read to be so limited. This is why extrinsic evidence of actual use by the applicant and registrant is essentially irrelevant in an ex parte likelihood of confusion analysis. The registrant's goods must be presumed to be sold to anyone, home user or hospital, through all normal trade channels, including AMAZON.COM as applicant sells its goods, and that home users and hospital supply procurement buyers are likely to encounter both applicant's and registrant's goods through these same trade channels.

Further, the registration(s) use(s) broad wording to describe its goods, namely, chemical reagents for medical diagnostic purposes, which presumably encompasses all goods and/or services of the type described, including applicant's more narrow identification of goods. In this case applicant's goods are diagnostic testing kits which consist of chemical reagents and assays used for testing for the presence of drugs, chemicals, hormones, viruses and bacteria to determine if the person tested has certain medical or health conditions. Applicant has since added a limitation that the goods are for home use only. The registrant's identification is not restricted as to any particular use or restricted to testing for any particular health condition. Therefore, it must be presumed to include all health conditions including those listed by the applicant such as drug intoxication, pregnancy, cancer, Covid-19, flu or genetic conditions. See, e.g., *Made in Nature, LLC v. Pharmavite LLC*, 2022 USPQ2d 557, at *44 (TTAB 2022); *In re Solid State Design Inc.*, 125 USPQ2d 1409, 1412-15 (TTAB 2018); *Sw. Mgmt., Inc. v. Ocinomled, Ltd.*, 115 USPQ2d 1007, 1025 (TTAB 2015). Thus, applicant's and registrant's goods are legally identical. See, e.g., *In re i.am.symbolic, llc*, 127 USPQ2d 1627, 1629 (TTAB 2018) (citing *Tuxedo Monopoly, Inc. v. Gen. Mills Fun Grp., Inc.*, 648 F.2d 1335, 1336, 209 USPQ 986, 988 (C.C.P.A. 1981); *Inter IKEA Sys. B.V. v. Akea, LLC*, 110 USPQ2d 1734, 1745 (TTAB 2014); *Baseball Am. Inc. v. Powerplay Sports Ltd.*, 71 USPQ2d 1844, 1847 n.9 (TTAB 2004)).

The attached evidence from BIO-TECHNE.COM, MDMAXX.COM, TEKNOVA.COM and AMAZON.COM show that goods such as chemical reagents for testing for medical conditions and testing kits which include medical reagents are often sold through the same retail channels to the same potential consumers. The fact that all of these websites feature "shopping carts" for easy ordering means that the ordinary home consumer and the medical lab technician can purchase these goods through the same website. Even a popular "sells all" consumer shopping website, AMAZON.COM, sells both reagents and test kits to anyone with money to buy them and for the specific purpose of testing for medical conditions such as COVID-19.

As explained above, the goods of the registrant have no restrictions as to nature, type, channels of trade, or classes of purchasers and are "presumed to travel in the same channels of trade to the same class of purchasers." *In re Vittera Inc.*, 671 F.3d 1358, 1362, 101 USPQ2d 1905, 1908 (Fed. Cir. 2012) (quoting *Hewlett-Packard Co. v. Packard Press, Inc.*, 281 F.3d 1261, 1268, 62 USPQ2d 1001, 1005 (Fed. Cir. 2002)); *Made in Nature, LLC v. Pharmavite LLC*, 2022 USPQ2d 557, at *49. Thus, applicant's and registrant's goods and/or services are related.

Because the goods of the parties are closely related, coupled with the fact that the marks of the parties are similar in their overall commercial impressions, consumers who encounter the marks SIGNIFY and SIGNIFY ANALYTICS could likely confuse the source of the goods.

DISCLAIMER REQUIREMENT - CONTINUED TO BE MADE FINAL

Applicant has argued that the disclaimer in this case is not required because the term "analytics" is vague and when coupled with the initial term "SIGNIFY" creates a unitary mark. However, applicant's mark is not unitary. A unitary mark would be one in which the wording is connected together to form a single word, e.g., SIGNIFYANALYTICS, in which case the disclaimer of the descriptive portion would not be required under office practice. A second means to form a unitary mark would be one which creates a bizarre incongruity or double entendre. Despite applicant's assertions to the contrary, there is nothing bizarre or incongruous about SIGNIFY ANALYTICS such as the marks DR. GRAMMAR or URBAN SAFARI create. Nor does it form a unitary slogan such as a mark like SIGNIFY YOUR COMMITMENT TO VITAL HEALTH ANALYTICS.

In this case the marks consists of separate words, SIGNIFY and ANALYTICS. The term "analytics" describes a scientific method for analyzing data. Applicant's goods are chemical reagent test kits which are designed to test for certain chemicals, hormones, viruses or drugs in the body. The kits are intended to collect data on what is contained in the bodily fluids tested and provide a means for the tester, be it a medical professional or a home user, to analyze the data collected by the test and determine why the person tested is sick, drunk, high, pregnant or whether two persons may be genetically related. The term, therefore, merely describes a key feature, characteristic or purpose of the goods.

Please call or email the assigned trademark examining attorney with questions about this Office action. Although an examining attorney cannot provide legal advice, the examining attorney can provide additional explanation about the refusal(s) and/or requirement(s) in this Office action. *See* TMEP §§705.02, 709.06.

The USPTO does not accept emails as responses to Office actions; however, emails can be used for informal communications and are included in the application record. *See* 37 C.F.R. §§2.62(c), 2.191; TMEP §§304.01-.02, 709.04-.05.

If applicant has already filed an appeal with the Trademark Trial and Appeal Board, the Board will be notified to resume the appeal. *See* TMEP §715.04(a).

If applicant has not filed an appeal and time remains in the response period for the final Office action, applicant has the remainder of that time to (1) [file another request for reconsideration](#) that complies with and/or overcomes any outstanding final requirement(s) and/or refusal(s), and/or (2) [file a notice of appeal](#) to the Board. TMEP §715.03(a)(ii)(B).

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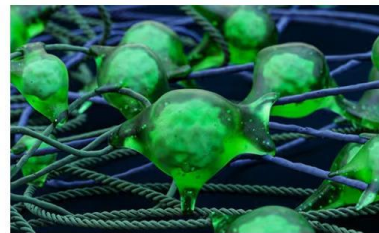
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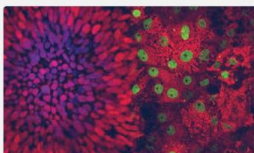
- Reduce variability in media composition
- Simplify compliance with regulatory guidelines
- Simplify comparability testing for raw material changes

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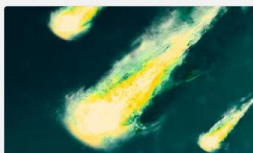
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Related Products for Cell Culture



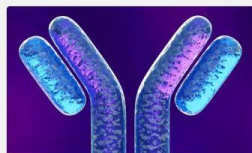
Differentiation Media, Supplements, and Kits

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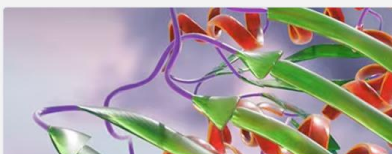
Cell Culture Assays

- Proliferation and viability assays
- Apoptosis assays
- CometAssay™ for DNA damage
- Invasion and migration assays
- Angiogenesis assay



Cell Isolation and Identification

- MagCelect™ Cell Isolation Kits
- Functional Identification Kits
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Cytokines and Growth Factors

- RUO, animal-free, and GMP grades
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Small Molecules and Peptides

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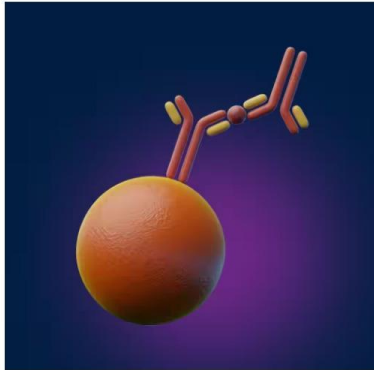
Our **Custom Cell Culture Media Manufacturing and Services** will work with you to expedite and standardize media production, develop optimized media formulations, and perform specialty media testing.

differentiation kits, and characterization kits for these specialty cell culture applications:

- Stem cell culture
- Organoid and 3D cell culture
- Immune cell culture
- Neural cell culture

Our expertise includes media and supplement production, media formula optimization, custom labeling, and assays for testing media on stem cells, immune cells, and other cell lines.

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From discovery to validation, you can trust our immunoassays to deliver precise, accurate results the first time and every time. Each of our immunoassays is based on our carefully tested, in-house antibodies, proteins, and specialized diluents.

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- ELISpot Kits

Validation

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- Immune Cell Protocols
- Neural Cell Culturing Protocols

Handbooks and Guides

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- The Organoid Culture Handbook
- Evolution of Cell Culture Model Systems eBook
- Flow Cytometry Handbook

Research Areas

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- Cell and Gene Therapy
- Transitioning to Animal-Free and GMP

Background Information

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Bio-Techne Brands

Which Brands are Currently Available on [bio-techne.com](#)? R&D Systems, Tocris Bioscience and ProteinSimple branded products are available to purchase through [bio-techne.com](#). ProteinSimple branded instruments are available to quote. ACD branded products will be available on [bio-techne.com](#) in the near future. Novus Biologicals branded products are not currently available on [bio-techne.com](#) and can be found at [novusbio.com](#).

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- ☐ Rat (6)
- ☐ Multi-Species (5)

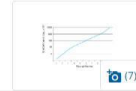
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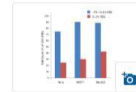
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
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GMP N-2 MAX Media Supplement (100X), Animal-free GMP

Catalog #: AR016

GMP Animal-free Media Supplement for Ex Vivo Cell or Tissue Manufacturing

Compare (Max 4): ☐



 (4)

View pricing ▾

StemXVivo Mesenchymal Stem Cell Expansion Media Best Seller

Catalog #: CCM004

★★★★★ (4)

Citations (12)

Compare (Max 4): ☐



 (3)

View pricing ▾

N-2 Plus Media Supplement

Catalog #: AR003

★★★★★ (2)

Citations (24)

Compare (Max 4): ☐



 (2)

View pricing ▾


ExCellerate B Cell Media, Xeno-Free Best Seller


Catalog #: CCM031

★★★★★ (1)

Expansion and clonal growth of B cells.

Compare (Max 4): ☐



 (5)

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QuantideX Portfolio of Assays with High Sensitivity and Streamlined Workflow



QuantideX qPCR BCR-ABL IS Kit

The first FDA-cleared Chronic Myeloid Leukemia (CML) monitoring assay provides deep sensitivity exceeding the NCCN guidelines with a Limit of Detection of MR4.7. Its multiplex design reduces hands-on time and increases throughput for labs.

[Learn More about the QuantideX qPCR BCR-ABL IS Kit](#)



QuantideX qPCR BCR-ABL Minor Kit*

With a workflow similar to the BCR-ABL IS kit, this assay allows labs to easily implement both assays to provide a more comprehensive testing solution.

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Recommended Accessories (3)

The image displays three product cards for Rhinowear prosthetics. Each card features a circular image of the product, a title, a price, a star rating, and a 'BUY NOW' button. The first card shows a young girl wearing a prosthetic lip, titled 'RhinoSwab Junior "Lips" Comfortable Self-Adhesive Silicone Lip Prosthetic' for \$7.25 with a 5-star rating. The second card shows a black prosthetic mustache, titled 'RhinoSwab Junior "Moustache" Comfortable Self-Adhesive Silicone Moustache' for \$7.25 with a 5-star rating. The third card shows a clear prosthetic nose, titled 'RhinoSwab Adult Comfortable Self-Adhesive Silicone Nose Prosthetic' for \$7.25 with a 5-star rating.

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Description

BinaxNOW COVID-19 Antigen Self Test (2 Tests): A simple solution for COVID-19 infection detection, with rapid results in the convenience of your home. This test has received FDA Emergency Use Authorization for self-testing without the need to ship samples to a lab or for a prescription from your healthcare provider. This 15-minute test can be completed anytime, anywhere.

Contact your Dedicated Account Manager

What's In The Box:

- (2) Sealed Test Cards
- (2) Foam Tipped Swabs
- (2) Dropper Bottles
- Illustrated Instructions

Manufacturer's Downloads



Guide

1.7 MB



User Manual

1.7 MB

Features

- Simple and easy to use, The BinaxNOW Self Test from Abbott requires just a shallow nasal swab that you can do yourself; includes easy-to-follow instructions.
- See results in 15 minutes, Convenient, fast results anytime, anywhere; no need for a prescription or sending to a lab.
- For ages 2 to adult, The self-test is indicated for children as young as 2 years old when administered by an adult, and for all people 15 and older to self-administer.
- Designed to detect active COVID-19 infection, Includes 2 tests that are indicated for serial testing—simply test yourself twice within 3 days, at least 36 hours apart; designed to detect active infection with or without symptoms.
- Does not include a virtual visit for certified digital test results. For proof of negative
- Detects active COVID-19 infection
- Includes illustrated step-by-step instructions
- Perform simple test procedure using a minimally invasive nasal swab
- Know your result now, without a need to wait for results from your healthcare provider
- It is available over the counter and does not require a prescription.
- The self-test is indicated for children as young as 2 years old when samples are collected by an adult, and for all people 15 and older.
- The test is for self-use when people test themselves twice within three days, with 36 hours between the two tests.

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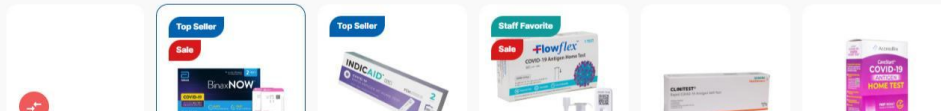
Count Per Pack:	2	Test Kit Dimensions:	9.125" L x 0.938" D x 5.063" H
Shipping Weight (in lbs):	0.22		

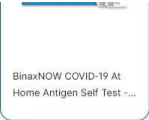
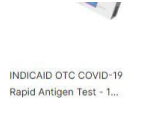

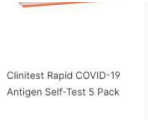

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Comparison Chart					
					
Price:	\$19.99 \$24.99	\$4.99	\$3.99 \$6.99	\$41.66	\$9.99
Manufacturer:	Abbott	PHASE Diagnostics	Acon	Siemens	AccessBio
Customer Rating:	★★★★★ 97 Reviews	★★★★☆	★★★★☆	★★★★☆	★★★★★ 2 Reviews

Frequently Asked Questions (FAQs)

What are expiration date?	+
Does it test for antibodies?	+
Can I use this test to meet travel requirements for testing?	+
What is the difference between an Antigen and Molecular test?	+
What is the performance of this test?	+
What do sensitivity and Specificity mean?	+
What ages is the test approved for use?	+
Should the test be stored and used at a certain temperature?	+
Does this test detect the new variants?	+

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Very good

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Reliable

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Very quick shipping!! Exactly as

Very quick shipping!! Exactly as expected.

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Lea S., Verified Buyer

01/10/22



BinaxNOW COVID-19 rapid test

This test was accurate and easy to use. Shipping was also fast and the test came well packaged and was handled with care. Would order again when I need more rapid tests!

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Nancy C. Verified Buyer

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★★★★★

Excellent service

I got my test kits even earlier than promised. The test was easy to use and I was grateful for the timely delivery.

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Lori P. Verified Buyer

01/06/22

★★★★★

Easy to use

Easy purchase; no issues

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Adelle I. Verified Buyer

01/06/22

★★★★★

Fast

Needed tests and received them quickly
Thank you! What a lifesaver.

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Delivered immediately as promised

The Covid antigen tests were promptly delivered as promised.

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Alice G. Verified Buyer

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prompt delivery

I had not heard of this company before, but I was looking for the Bina tests. I took a chance, and the delivery was prompt, items are in good condition. I only wish everything was not all sold out now.

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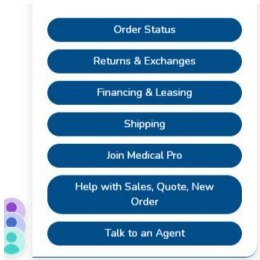
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Description

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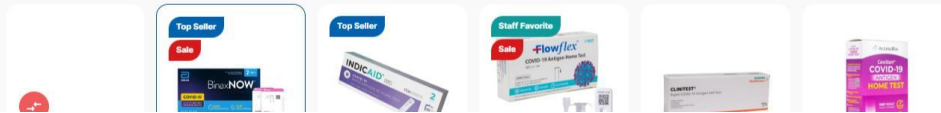
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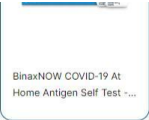
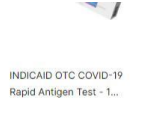

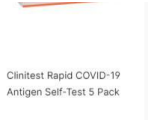

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Very good

Very good

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Reliable

Very reliable

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- Excellent service**
 I got my test kits even earlier than promised. The test was easy to use and I was grateful for the timely delivery.

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Easy to use
 Easy purchase; no issues

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Fast
 Needed tests and received them quickly
 Thank you! What a lifesaver.

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Delivered immediately as promised
 The Covid antigen tests were promptly delivered as promised.

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prompt delivery
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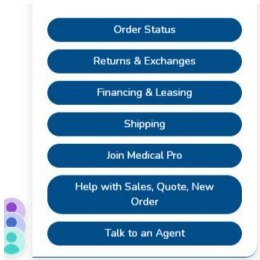
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Reagent i-STAT CG4+ Blood Gases Lactate For i-STAT...

\$350

MFG Name: Abbott
SKU: MD550532
MFG Part: 03P8551

☆☆☆☆☆

Reagent Architect Core Hepatitis / Antibody Test Anti...

\$1,135.18

MFG Name: Abbott
SKU: MD613683
MFG Part: 06L2225

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Reagent Architect Antibody Test Hepatitis B Surface Antigen Fo...

\$975.34

MFG Name: Abbott
SKU: MD613688
MFG Part: 04P5325

☆☆☆☆☆

Reagent i-STAT Chem8+ Cartridge

\$359

MFG Name: Abbott
SKU: MD550531
MFG Part: 09P31-26

☆☆☆☆☆

☐ Germany

☐ USA



Clarify 10SG Urine Reagent Strips

\$16.92

MFG Name: **Clarity Diagnostics**
SKU: **MD475760**
MFG Part: **CLA-URS10**
☆☆☆☆☆



Ketostix Reagent Ketone Test Strips for Urinalysis

\$384.34

MFG Name: **Ascensia Diabetes Care**
SKU: **MD438310**
MFG Part: **2880**
☆☆☆☆☆



Urinalysis Test Strips, CLIA Waived

\$423.38

MFG Name: **Ascensia Diabetes Care**
SKU: **MD438309**
MFG Part: **2882**
☆☆☆☆☆



Urocheck Urine Reagent Strips 2GP

\$15.38

MFG Name: **Clarity Diagnostics**
SKU: **MD475765**
MFG Part: **DTG-2GP**
☆☆☆☆☆



Ketostix Reagent Ketone Test Strips for Urinalysis

\$345.71

MFG Name: **Ascensia Diabetes Care**
SKU: **MD438311**
MFG Part: **2881**
☆☆☆☆☆

 Diastix Reagent Strips

Diastix Reagent Strips

\$240.57

MFG Name: **Ascensia Diabetes Care**
SKU: **MD438307**
MFG Part: **2803**
☆☆☆☆☆



Urinalysis Test Strips, CLIA Waived

\$464.09

MFG Name: **Ascensia Diabetes Care**
SKU: **MD438306**
MFG Part: **2883**
☆☆☆☆☆



Clarity Urocheck 7 Urine Test Strips

\$30.85

MFG Name: **Clarity Diagnostics**
SKU: **MD475762**
MFG Part: **DTG-URO7**
☆☆☆☆☆

Sale



Consult 10SG Urine Reagent Strips

\$35

\$44.58

MFG Name: Mckesson
SKU: MD58876
MFG Part: 121-10SG

☆☆☆☆☆

Urocheck Urinalysis Reagent Test Strips 5 OB

\$19.3

MFG Name: Clarity Diagnostics
SKU: MD475763
MFG Part: DTG-50B

☆☆☆☆☆

Diastix Reagent Strips

\$271.26

MFG Name: Ascensia Diabetes Care
SKU: MD438308
MFG Part: 2806

☆☆☆☆☆


Pentra 400 Reagent Container

\$51.51

MFG Name: Globe Scientific
SKU: MD471653
MFG Part: 6507

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Reagent Cholestech LDX Lipid Profile & Glucose Cassette For...

\$143.13

MFG Name: Abbott
SKU: MD818298
MFG Part: 10-991

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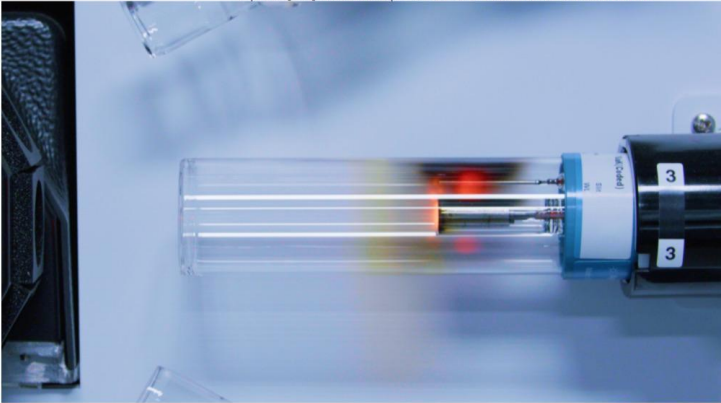
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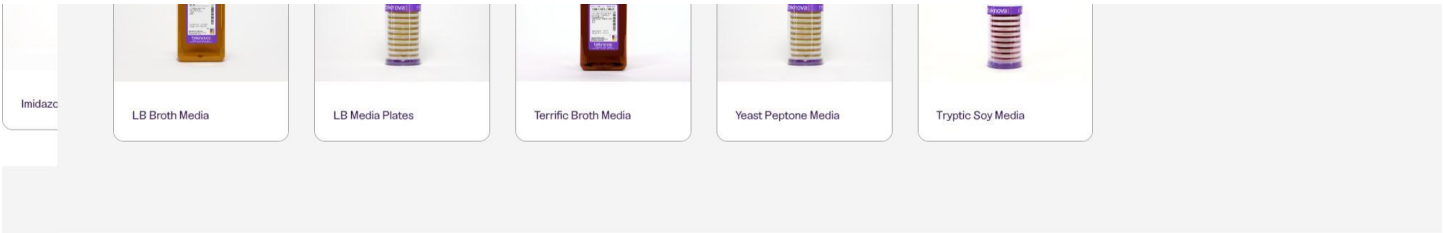
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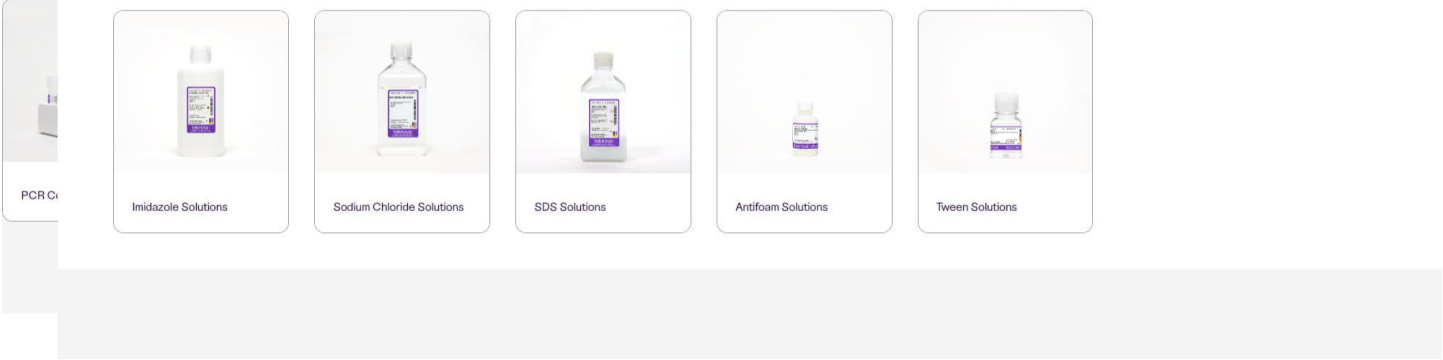
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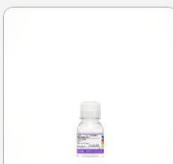
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PCR Certified Water



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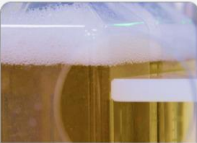
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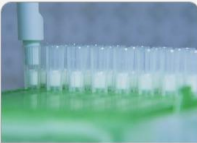
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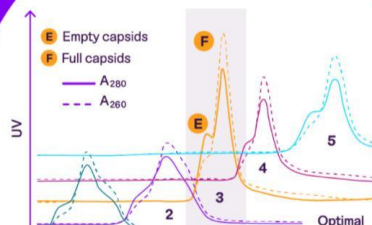
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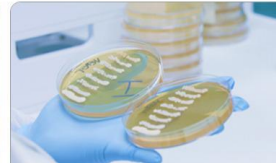
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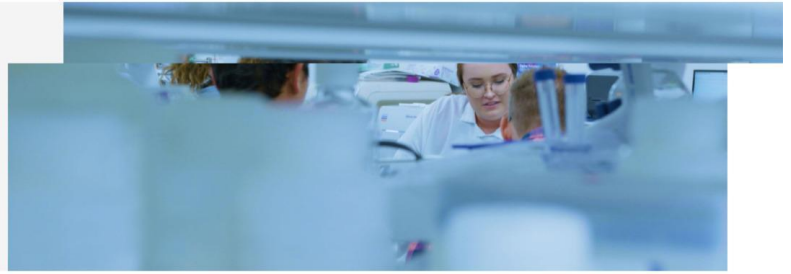
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110 Piece Set

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30 Count (Pack of 1)


★★★★☆ ~ 205

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1 Count (Pack of 1)

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